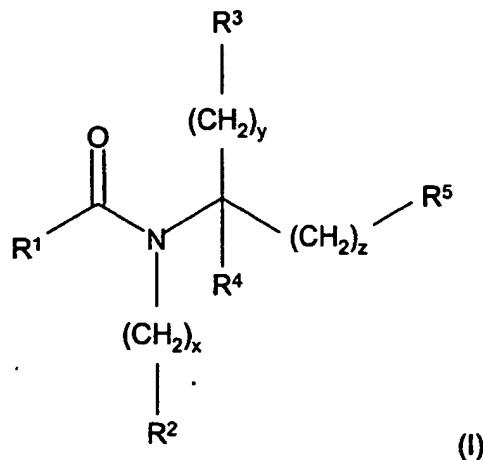


IN THE CLAIMS

1. (canceled).

2. (Previously presented) A compound of formula (I)



wherein:

R^1 is selected from:

- a) phenyl, which is optionally substituted by 1-3 groups each independently selected from C_1-C_6 alkyl, CF_3 , halo, CN , NR^7R^8 , SO_2R^6 and OC_1-C_6 alkyl, and
- b) Aromatic Heterocycle, wherein said Aromatic Heterocycle is selected from pyridyl, pyrazinyl, pyrimidinyl, quinolinyl, quinoxalinyl, isoxazolyl and pyrazolyl, each aromatic heterocycle optionally substituted by 1-3 groups each independently selected from C_1-C_6 alkyl, SR^6 , SO_2R^6 , NH_2 , CF_3 , halo, OH , OC_1-C_6 alkyl, NR^7R^8 wherein R^8 may be optionally substituted by NH_2 , phenyl or Heterocycle, and OPh wherein Ph may be optionally substituted by 1-3 groups each independently selected from halo and C_1-C_6 alkyl;

R^2 is selected from:

- a) phenyl, which is optionally substituted by methyl, fluoro, chloro, methoxy, CF_3 or SO_2CH_3 ,
- b) pyrazolyl, which is optionally substituted by methyl, and
- c) $C(O)N(CH_3)_2$;

R^3 is selected from:

- a) phenyl, said phenyl being optionally fused to Heterocycle and said phenyl or said fused phenyl being optionally substituted by 1-3 groups each independently selected from C₁-C₆ alkyl, halo, CN and OC₁-C₆ alkyl,
- b) R⁶,
- c) cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl, which is optionally substituted by C₁-C₆ alkyl; and
- d) Aromatic Heterocycle, wherein said Aromatic Heterocycle may be defined as a 5-6 membered aromatic heterocycle containing 1 or 2 nitrogen atoms, said ring optionally fused with a phenyl or a 3-8 membered cycloalkyl group.

R⁴ is H;

R⁵ is CONH₂;

R⁶ is methyl;

R⁷ is hydrogen or C₁-C₆ alkyl;

R⁸ is C₁-C₆ alkyl;

or NR⁷R⁸ forms a monocyclic saturated ring system containing between 3 and 7 ring atoms;

x is 1;

y is 0; and

z is 0 or 1

wherein:

Aromatic Heterocycle may be defined as a 5-6 membered aromatic heterocycle containing 1-4 heteroatoms each independently selected from N, O and S, said ring optionally fused with a phenyl or a 3-8 membered cycloalkyl group;

Heterocycle is a 5-8 membered saturated or partially saturated ring containing 1-3 heteroatoms each independently selected from N, O and S, said ring optionally fused with phenyl;

a tautomer thereof or a pharmaceutically acceptable salt, solvate or polymorph of said compound or tautomer.

2 3. (Currently amended) A compound according to claim 2 wherein R¹ is selected from:

- a) phenyl, which is optionally substituted by 1-3 groups each independently selected from C₁-C₆ alkyl, CF₃, halo, CN, NR⁷R⁸, SO₂R⁶ and OC₁-C₆ alkyl, and
- b) Aromatic Heterocycle, wherein said Aromatic Heterocycle is selected from:
 - i) pyridyl, which is optionally substituted by 1-3 groups each independently selected from C₁-C₆ alkyl, SO₂R⁶, NH₂, CF₃, CN, halo, OH, OC₁-C₆ alkyl, NR⁷R⁸ wherein R⁸ may be optionally substituted by NH₂, phenyl or Heterocycle, and OPh wherein Ph may be optionally substituted by 1-3 groups each independently selected from halo and C₁-C₆ alkyl;
 - ii) pyrimidinyl, which is optionally substituted by 1-3 groups each independently selected from C₁-C₆ alkyl, SO₂R⁶, NH₂, CF₃, CN, halo, OH, OC₁-C₆ alkyl, NR⁷R⁸ wherein R⁸ may be optionally substituted by NH₂, phenyl or Heterocycle, and OPh wherein Ph may be optionally substituted by 1-3 groups each independently selected from halo and C₁-C₆ alkyl;
 - iii) pyrazinyl, which is optionally substituted by 1-3 groups each independently selected from C₁-C₆ alkyl, NH₂, SR⁶ and halo;
 - iv) quinolinyl;
 - v) quinoxalinyl, which is optionally substituted by OH;
 - vi) isoxazolyl, which is optionally substituted by 1-3 groups each independently selected from: C₁-C₆ alkyl; and
 - vii) pyrazole;

R² is selected from:

- a) phenyl, which is optionally substituted by methyl, halo, methoxy, CF₃ or SO₂CH₃;
- b) cyclopropyl or 1- or 2-indanyl;
- c) pyrazolyl, which is optionally substituted by methyl;
- d) C(O)N(CH₃)₂, and
- e) piperidinyl substituted by C(O)R⁶.

R^3 is selected from:

- a) phenyl, said phenyl being optionally fused to 1,4-dioxan and said phenyl or said fused phenyl being optionally substituted by 1-3 groups each independently selected from $C_1.C_6$ alkyl, halo, CN and $OC_1.C_6$ alkyl;
- b) R^6 ,
- c) cyclopropyl, which is optionally substituted by $C_1.C_6$ alkyl; and
- d) Aromatic Heterocycle, wherein said Aromatic Heterocycle is selected from pyrazolyl or pyridyl, both optionally substituted by $C_1.C_6$ alkyl;

R^5 is $CONH_2$ or CH_3 ; and

z is 0.

³
²
⁴ A. (Currently amended) A compound according to any one of claims 1 to ~~2~~ or ~~3~~ wherein R^1 is phenyl, 2- or 3-pyridyl or 2,4-pyrimidinyl, said moieties being optionally substituted by 1-3 groups each independently selected from $C_1.C_6$ alkyl, halo, $OC_1.C_6$ alkyl, CN, SO_2R^6 , NHR_7 , $NHCH_2CH_2NH_2$ and CF_3 ;

⁴
⁵ 5. (original) A compound according to claim ⁴ wherein R^1 is phenyl, 2- or 3-pyridyl or 2,4-pyrimidinyl, said moieties being optionally substituted by 1-3 groups each independently selected from methyl, fluoro, chloro, methoxy, ethoxy, n-propoxy, CN, SO_2CH_3 , NH_2 , $NHCH_3$, $NHCH_2CH_2NH_2$, and CF_3 .

6. (canceled)

⁵
⁷ 7. (previously presented) A compound according to claim ⁵ wherein R^2 is phenyl, para-fluorophenyl, para-chlorophenyl, para-methylphenyl, 2,5-dimethylphenyl, o-methylphenyl and para-methoxyphenyl.

⁶
⁸ 8. (previously presented) A compound according to claim ⁷ wherein R^3 is selected from:

- a) phenyl, said phenyl being optionally fused to 1,4-dioxan and said phenyl or said fused phenyl being optionally substituted by 1-2 groups each independently selected from methyl, methoxy, ethoxy, fluoro, chloro and CN;

- b) isopropyl;
- c) cyclopropyl; and
- d) pyrazolyl and pyridyl, both optionally substituted by methyl.

6

7 9. (original) A compound according to claim 8 wherein R³ is 3-methoxyphenyl or 1,4-benzodioxanyl.

10. (Cancelled).

)

8 11. (Previously presented) A compound according to claim 2 selected from:

2-Amino-N-[2-amino-1-(2-methylphenyl)-2-oxoethyl]-N-(4-chlorobenzyl)nicotinamide,

N-[2-Amino-1-(3-methoxyphenyl)-2-oxoethyl]-4-cyano-N-(4-methylbenzyl)benzamide,

N-[3-Amino-1-(3-methoxyphenyl)-3-oxopropyl]-4-methyl-N-(4-methylbenzyl)nicotinamide,

2-Amino-N-[(1S)-3-amino-3-oxo-1-phenylpropyl]-N-(4-methylbenzyl)nicotinamide,

5-Chloro-2-methylthio-N-[2-amino-1-{1,4-benzodioxan-6-yl}-2-oxoethyl]-N-(4-methylbenzyl)pyrimidine-4-carboxamide,

5-Chloro-2-amino-N-[2-amino-1-{1,4-benzodioxan-6-yl}-2-oxoethyl]-N-(4-methylbenzyl)pyrimidine-4-carboxamide, and

2-Amino-N-[carbamoyl-(2,3-dihydro-benzo[1,4]dioxin-6-yl)-methyl]-4,6-dimethyl-N-(4-methyl-benzyl)-nicotinamide;

and tautomers thereof and pharmaceutically acceptable salts, solvates and polymorphs of said compound or tautomer.

9 12. (Previously presented) A pharmaceutical composition comprising a compound of claim 2, or pharmaceutically acceptable salts, solvates or polymorphs thereof, and a pharmaceutically acceptable diluent or carrier.

10 13. (previously canceled)

10 14. (Previously presented) A method of treatment of a disorder or condition where inhibition of Oxytocin is known, or can be shown, to produce a beneficial